



SERIOUS EVENT REPORTING

Please fax or email a copy of this document as soon as possible after a serious adverse event is detected.

Fax : 514-832-0266

Email : HQ-cases.managers@hema-quebec.qc.ca

Event type

- Adverse event – Recipient
 Adverse event – Donor
 Accident / Error – Product

Product type

- Cord Blood Unit
 Bone Marrow
 Peripheral Blood Stem Cells

Recipient ID: _____

Donor ID: _____

Transplant Date (dd/mm/yyyy) : _____

Date the event was detected (dd/mm/yyyy) : _____

Recipient file ID: _____

Hospital Name: _____ Code: _____

Physician Name: _____

Phone number: _____ ext: _____

Faxed on (dd/mm/yyyy) : _____ By: _____ S/O

Description of the event

Signature (physician or representative)

Date: _____
(dd/mm/yyyy)